

INFORMATION DISCLOSURE CITATION			Docket No.: RLL-283US		Serial No.: 10/540,062	
			Applicants: M [REDACTED] TA et al.			
			Filed: 6/22/2005		Group: 1625	

U.S. PATENT DOCUMENTS							
EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
YD	A1	3,176,019	3/30/1965	Campbell et al.	260	293.4	
	A2	5,281,601	1/25/1994	Cross et al.	514	320	
	A3	5,948,792	9/7/1999	Tsuchiya et al.	514	317	
	A4	6,130,232	10/10/2000	Mase et al.	514	318	
↓	A5	6,174,900	1/16/2001	Okada et al.	514	317	

FOREIGN PATENT DOCUMENTS

		DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUBCLASS	TRANSLATION YES NO
YD	B1	EP 0 072 620	2/23/1983	EPO	C07D	405/12	1
	B2	EP 0 108 986	5/23/1984	EPO	C07D	311/30	1
	B3	EP 0 267 319	5/18/1988	EPO	C07C	153/11	1
	B4	EP 0 325 571	7/26/1989	EPO	C07C	215/54	1
	B5	EP 0 388 054	9/19/1990	EPO	C07D	207/08	1
	B6	EP 0 801 067	10/15/1997	EPO	C07D	453/02	1
	B7	GB 940,540	10/30/1963	UK	C07C		1
	B8	JP 135958/1994	5/17/1994	Japan	C07D	333/16	1
	B9	JP 92921/1994	4/5/1994	Japan	C07C	237/20	1
	B10	WO 91/09013	6/27/1991	PCT	C07D	207/08	1
	B11	WO 93/16018	8/19/1993	PCT	C05F	17/02	1
	B12	WO 93/16048	8/19/1993	PCT	C07D	211/26	Equivalent CA 2155320
	B13	WO 96/33973	10/31/1996	PCT	C07D	211/46	Equivalent US 5,750,540
	B14	WO 97/45414	12/4/1997	PCT	C07D	211/58	Abstract
	B15	WO 98/05641	2/12/1998	PCT	C07D	211/46	Equivalent US 5,948,792
↓	B16	WO 98/29402	7/9/1998	PCT	C07D	311/20	1

EXAMINER	T-A. Sola	DATE CONSIDERED	9-11-07
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*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)		
Yb	C1	Kubo et al., "Cloning, sequencing and expression of complementary DNA encoding the muscarinic acetylcholine receptor", <i>Nature</i> , 323(2):411-416 (1986)
	C2	Bonner et al., "Identification of a Family of Muscarinic Acetylcholine Receptor Genes", <i>Science</i> , 237:527-531 (1987)
	C3	Eglen et al., "Muscarinic receptor ligands and their therapeutic potential", <i>Current Opinion in Chemical Biology</i> , 3:426-432 (1999)
	C4	Eglen et al., "Therapeutic opportunities from muscarinic receptor research", <i>Trends in Pharmacological Sciences</i> , 22(8):409-414 (2001)
	C5	Felder et al., "Therapeutic Opportunities for Muscarinic Receptors in the Central Nervous System", <i>Journal of Medicinal Chemistry</i> , 43(23):4333-4353 (2000)
	C6	Broadley and Kelly, "Muscarinic Receptor Agonists and Antagonists", <i>Molecules</i> , 6:142-193 (2001)
	C7	Birdsall et al., "Muscarinic receptors: it's a knockout", <i>Trends in Pharmacological Sciences</i> , 22(5):215-219 (2001)
	C8	de Groat and Yoshimura, "Pharmacology of the Lower Urinary Tract", <i>Annual Review of Pharmacology and Toxicology</i> , 41:691-721 (2001)
	C9	Steers, "The future direction of neuro-urology drug research", <i>Current Opinion in CPNS Investigational Drugs</i> , 2(3):268-282
	C10	Chapple, "Muscarinic receptor antagonists in the treatment of overactive bladder", <i>Urology</i> , 55(Suppl. 5A):33-46 (2000)
	C11	Steers, Barrot, Wein, "Voiding dysfunction: diagnosis classification and management", In: <i>Adult and Pediatric Urology</i> , ed. Gillenwater, Grayhack, Howards, Duckett. Mosby, St. Louis, MO; 1220-1325, 3rd edition (1996)
	C12	Sagara et al., "Cyclohexylmethylpiperidinyltrifluoromethylpropioamide: A Selective Muscarinic M ₃ Antagonist Discriminating against the Other Receptor Subtypes", <i>Journal of Medicinal Chemistry</i> , 45:984-987 (2002)
	C13	Braish et al., "Construction of the (1 α ,5 α ,6 α)-6-Amino-3-azabicyclo[3.1.0]hexane Ring System", <i>Synlett</i> , 1100-1102 (1996)
	C14	Moriya et al., "Affinity Profiles of Various Muscarinic Antagonists for Cloned Human Muscarinic Acetylcholine Receptor (mAChR) Subtypes and mAChRs in Rat Heart and Submandibular Gland", <i>Life Sciences</i> , 64(25):2351-2358 (1999)
↓	C15	Cheng and Prusoff, "Relationship between the inhibition constant (K _i) and the concentration of inhibitor which causes 50 per cent inhibition (I ₅₀) of an enzymatic reaction", <i>Biochemical Pharmacology</i> , 22:3099-3108 (1973)

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